This estimate is based on the average number of new color additive petitions received in 1997 and 1998. Although the burden varies with the type of petition submitted, a color additive petition involves analytical work and appropriate toxicology studies, as well as the work of drafting the petition itself. Because labeling requirements under § 70.25 for a particular color additive involve information required as part of the color additive petition safety review process, the estimate for the number of respondents is the same for § 70.25 as for § 71.1, and the burden hours for labeling are included in the estimate for § 71.1.

Color additives are subjected to payment of fees for the petitioning process. The listing fee for a color additive petition ranges from $1,600 to $3,000, depending on the intended use of the color and the scope of the requested amendment. A complete schedule of fees is set forth in 21 CFR 70.19. An average of two Category A and three Category B color additive petitions are expected per year. The maximum color additive petition fee for a Category A petition is $2,600 and the maximum color additive petition fee for a Category B petition is $3,000. Because an average of five color additive petitions are expected per calendar year, the estimated total annual cost burden to petitioners for this startup cost would be less than or equal to $14,200 (2 x $2,600 + 3 x $3,000 listing fees = $14,200).

Dated: April 5, 1999.

William K. Hubbard,

Acting Deputy Commissioner for Policy.

[FR Doc. 99–8954 Filed 4–9–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99D–0297]

Draft Guidance for Industry on Formal Dispute Resolution; Appeals Above the Division Level; Availability; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of March 19, 1999 (64 FR 13587). The document announced an opportunity for public comment on a draft guidance entitled ''Formal Dispute Resolution; Appeals Above the Division Level.'' The notice was published with two inadvertent errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In FR Doc. 99–6749, appearing on page 13587 in the **Federal Register** of Friday, March 19, 1999, the following corrections are made:

1. On page 13587, in the first column, under the **DATES** caption, the last sentence is corrected to read ''Written comments on the information collection provisions must be submitted to the Dockets Management Branch (address below) by May 19, 1999. All comments should be identified with the docket number found in brackets in the heading of this document.''

2. On page 13589, in first column, the last paragraph of the document is removed.

Dated: April 6, 1999.

William K. Hubbard,

Acting Deputy Commissioner for Policy.

[FR Doc. 99–8997 Filed 4–9–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99D–0296]

Draft Guidance for Industry on Formal Meetings with Sponsors and Applicants for PDUFA Products; Availability; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of March 19, 1999 (64 FR 13591). The document announced an opportunity for public comment on a draft guidance entitled ''Formal Meetings with Sponsors and Applicants for PDUFA Products.'' The notice was published with two inadvertent errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In FR Doc. 99–6748, appearing on page 13591 in the **Federal Register** of Friday, March 19, 1999, the following corrections are made:

1. On page 13591, in the third column, under the **DATES** caption, after the last sentence two sentences are added to read ''Written comments on the information collection provisions must be submitted to the Dockets Management Branch (address below) by May 19, 1999. All comments should be identified with the docket number found in brackets in the heading of this document.''

2. On page 13594, in the first column, the last paragraph of the document is removed.

Dated: April 6, 1999.

William K. Hubbard,

Acting Deputy Commissioner for Policy.

[FR Doc. 99–8998 Filed 4–9–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–460]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Extension of a currently approved collection;

Title of Information Collection: Medicare Participating Physician or Supplier Agreement, HCFA–460;

Form No.: HCFA–460 (OMB# 0938–0373);

Use: The HCFA–460 is completed by nonparticipating physicians and